



Development of Durable Materials for cost Effective AWS Utilizing All-Ceramic Solid Oxide Electrolyzer Stack Technology

Dr. John Pietras Saint-Gobain 5/30/2020





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Project Overview

Project Partners

Dr. John Pietras, Saint-Gobain Dr. Srikanth Gopalan, Boston University Dr. Jeffry Stevenson, PNNL Dr. Olga Marina, PNNL

Project Vision

Novel chemistries of nickelate-based materials showing enhanced oxygen hyperstoichiometry are being developed to solve the issue of air electrode delamination during SOEC operation

Project Impact

The steady state degradation rate of SOEC stacks will be improved by solving the issue of electrode delamination. The materials developed will be compatible with the highly stable (0.2%/khr degradation rate) co-sintered SOFC stack architecture and result in a cost effective H_2 production platform

* this amount does not cover support for HydroGEN resources leveraged by the project (which is provided separately by DOE) HydroGEN: Advanced Water Splitting Materials

ward #	EE0008377
tart/End Date	10/01/2018 – 9/31/2021
ear 1 Funding*	\$0.312M
ost Share %	20%





Approach: Summary

Project Motivation

A previous project at Boston University funded by Saint-Gobain showed that Lanthanum nickelate – GDC composites could improve SOFC performance.

A solution to the decomposition issue when LNO is in contact with GDC was developed.

This project was initiated to extend this research to SOEC performance and investigate more promising material sets

Barriers

Phase stability/performance (Boston University)

Identification of phase stability boundaries with target electrochemical properties

Co-sintering (Saint-Gobain)

Incorporate materials within stacks ensuring porosity, activity, defect free microstructure

Accelerated testing (PNNL)

Development of a protocol which probes the dominate degradation mechanism

Key Impact

Metric	State of the Art	Expected Advance
ASR	0.3-0.5 ohm cm ²	<u><</u> 0.3 ohm cm ²
Current Density	0.5 A/cm ²	<u>></u> 1 A/cm² @ 1.4V
Degradation Rate	1-4 %/khr	<u><</u> 0.3 %/khr

Partnerships

Saint Gobain (Dr. J. Pietras) provides an expertise in materials development and extensive US manufacturing footprint. Has developed an extremely stable all-ceramic, co-fired SOFC solution with a degradation rate of 0.2%/khr

Boston University (Dr. S. Gopalan) has demonstrated stable nickelate chemistries and draws on expertise in advancing the chemistry of electrochemical devices

PNNL (Dr J. Stevenson) has developed in-situ characterization capabilities to monitor cells and electrochemical interfaces along with expertise in design and interpretation of acceleration testing

Approach: Problem Definition

Leverage SOFC Work

 Saint-Gobain has developed a unique cosintering process to produce SOFCs from button cells to stacks



Potential of Nickelate Family: Performance + Durability

- Rare-earth nickelates have an open alternating crystal structure and thus a large number of oxygen interstitial sites
- Oxygen exchange and transport is higher at higher oxygen pressures than in state-of-the-art perovskite oxides leading to high performance
- Voids and cracks can be avoided by incorporating oxygen in the open lattice



V.V. Kharton, et. al, J.Solid.State.Chem., 181, 1425-1433 (2008)

Issue to solve: Reaction with doped ceria

 Decomposition of nickelate phase when in contact with doped Ceria



Approach: Innovative Solution

Strategy to Stabilize LNO Phase

- Push to highly doped Ceria compositions
- Operate on the pseudo-binary tie-line between LNO and the saturated ceria phase
- Prevent incorporation of lanthanides into the ceria structure thus preserving nickelate phase



Project Approach

- Synthesize material compositions and utilize XRD to establish phase purity of end compounds
- Create ceria nickelate mixtures and measure stability & performance characteristics
- Sinter button cells utilizing both conventional and co-sintering techniques, optimizing microstructure
- Electrochemically test button cells for performance and durability

	2018			2019								
Calendar Period	10	11	12	1	2	3	4	5	6	7	8	9
Project Period	M1	M2	M3	M4	M5	M6	M7	M8	M9	M10	M11	M12
1.Materials Development and Transport Property Measurements												
1.1 Composite powder production												
1.2 Determine oxygen hyperstoichimetry												
1.3 Measurement of composite properties												
1.4 Point defect chemistry of composites												
M1.1: Phase pure powders (stoichiometry & XRD confirmation)			м									
M1.2: Hyperstoichiometry and identify phase stability boundaries for nickelates & doped												
ceria								М				
M1.3: ID top 3 candidates (conductivity >96 S/cm, oxygen exchange coefficient 10 ⁻⁵ cm/s)												М
2. Degradation Analyses on Materials and Interfaces												
2.1 Preparation of test rig												
2.2 Button cell fabrication, current technology												
2.3 Button cell testing & characterization, current technology												
M2.1: Baseline button cells, He leak <1x10 ⁻⁸ mbarL ⁻¹ cm ⁻²						М						
M2.2: Degradation mechanisms confirmed and accelerated testing developed											М	
Go/No-Go Decision Point (Phase I)												G1

Approach: Summary of Results

Developed, produced and tested advanced electrodes with improved performance over baseline material set





- The goal of the proposed effort is to develop a fundamental understanding of performance degradation and electrode delamination in nickelate based SOEC materials and interfaces, and to develop compositions addressing this degradation while meeting specific performance targets:
 - ASR ≤0.30 ohm-cm2
 - current density >1A/cm2 at 1.4V
 - stack electrical efficiency >95% LHV H2
 - stack lifetime ≥7 years
- High oxygen pressure at the electrolyte-electrode interface has been identified in the literature and in ongoing EERE funded projects as a major cause for degradation within SOEC. The build-up of oxygen pressure can be mitigated by incorporating materials with high oxygen hyper stoichiometries close to the electrolyte-anode interface.
- Thus this project with look to generate solutions based on previous Consortium modeling. In this way the program will push the state of the art and encourage focused discussion on the topic among the Consortium.

Accomplishments: Powder Synthesis



Neodymium System Example

- Synthesis of Neodymium Nickelate (Nd₂NiO₄) and Neodymium Doped Ceria (NDC) by solid state reactions
- 10-50 mol% neodymium was doped into the ceria
- XRD of the NDC powders confirms the fluorite crystal structure even in highly doped ceria
- Peak shift indicates the increase of the lattice parameter with increasing dopant concentration
- Replacement of smaller Ce⁴⁺ ions with the larger Nd³⁺ ions (the ionic radii of Ce⁴⁺ and Nd³⁺ are 0.97 and 1.1053 Å respectively) leads to the cubic ceria lattice expansion.







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Accomplishments: Stability in Nd System

50/50 MIXTURES OF $\rm Nd_2NiO_4$ WITH ND DOPED CERIA CALCINED AT 1300 °C CONFIRM THE STABILITY HYPOTHESIS



No nickelate phase retained

Nickelate phase retained



STABILITY HYPOTHESIS DUPLICATED IN THE La SYSTEM



Shift in lattice parameter from starting single phase value indicates incorporation of La into the ceria lattice for dopant concentrations <40 mol%



HIGH CTE OF LNO CAN BE REDUCED BY CERIA ADDITION BUT IS AN AREA OF CONCERN



High CTE of LNO can be reduced by ceria

addition but remains an area of concern

Coefficient of thermal expansion (1/K)						
LDC	13.2 × 10 ⁻⁶					
LNO	15.1 × 10 ⁻⁶					
NDC	14.4 × 10 ⁻⁶					
NNO	15.9 × 10⁻ ⁶					

Minimal chemical expansion on pO₂ change



Linear expansion as a function of time at 800°C showing a transition from 1 atm to 10^{-5} atm pO₂ for LNO, NNO, LDC and NDC. All of the materials show only a small change in dimensions.



Accomplishments: TGA of LaNiO_x & NdNiO_x

NICKELATES SHOW ABILITY TO STORE AND RELEASE OXYGEN, POSSIBLE GREATER AFFINITY FOR OXYGEN IN NO BASED NICKELATE THAN LA BASED

- As produced powders show very little uptake/release of oxygen
- Samples were pre-conditioned in Ar at 1000°C.
- TGA profile for LaNiO_x virtually independent of p(O₂).
- NdNiO_x shows greater O₂ uptake/release on gravimetric basis.
- Some change at 330°C which causes loss of oxygen with increasing temperature.





Accomplishments: Oxygen Uptake



Greater oxygen update for nickelates compared to LSM

Step-wise pO2 change confirms Nd has more open structure then La nickelate



Results confirmed through peak shift

Shift towards right indicates unit cell shrinkage under low pO₂





PROCEDURE DEVELOPED TO ISOLATE SURFACE EXCHANGE REACTION

Constant current of 1 A is applied and voltage is measured

$$\sigma = \frac{LI}{AV}$$

Porous bars are used to reduce the diffusion length

- Oxygen Diffusion no longer rate limiting
- Relaxation time, t = l²/D ~ 2 sec



 $I = diffusion \ length = 5 \ \mu m$ $D = 1.5 \ x 10^{-7} \ cm^2/s$ LNO @ 800°C

Surface exchange is the only rate limiting step

$$\sigma_n = \frac{\sigma(t) - \sigma(0)}{\sigma(\infty) - \sigma(0)} = 1 - \exp(-\frac{kt}{l})$$

Testing Sequence



Accomplishments: Conductivity Relaxation

NORMALIZED CONDUCTIVITY RESULTS SUGGEST NICKELATES WILL OUTPERFORM LSM

- Both LNO & NNO show 5 orders of magnitude improvement over LSM
- Transitions occur faster when going from low pO₂ to higher pO₂ aka in SOEC mode
- Faster oxygen transport kinetics in SOEC mode

	Exchange coefficient (m/s)			
	1-10%	10-1%		
LNO	1.40x10 ⁻⁶	5.66x10 ⁻⁷		
NNO	1.59x10 ⁻⁶	6.19x10 ⁻⁷		

LSM = 1x10⁻¹² (m/s) Phys. Chem. Chem. Phys. 2013, 15, 2298 Effect of A and B-site cations on surface exchange coefficient for ABO₃ perovskite materials Eric N. Armstrong,_{*} Keith L. Duncan, and Eric D. Wachsman*.







Accomplishments: Electrical Conductivity

CONDUCTIVITY MATCHES THE TARGET OF >96 S/cm

Conductivity variation as a function of vol% nickelate

- Percolation limit in these mixtures occurs at ~ 40 vol%
- Below nickelate percolation limit conductivity is controlled by ceria (ionic conductivity)
- Above nickelate percolation limit conductivity is controlled by nickelate (electronic conductivity)





Accomplishments: Symmetrical Cells

NICKELATE STRUCTURES SHOW MUCH LOWER POLARIZATION RESISTANCE THAN LSM BASED ELECTRODES





ABLE TO INCORPORATE NEW MATERIALS INTO THE CO-SINTERING PROCESS



Uniform baseline microstructure



Improved LNO/LDC electrode microstructure





STABILIZED NICKELATES SHOW IMPROVEMENT IN PERFORMANCE OVER BASELINE

- >80% improvement in current density
- Good agreement between 3 labs
- Performance improvement shown on:
 - Anode supported cells (BU cell)
 - Co-sintered cells (SG cell)
- Solution can be utilized industry wide



STABILIZED NICKELATES SHOW ABILITY TO FUNCTION IN SOEC AND SOFC MODES



Accomplishments: Durability Testing

NICKELATE SYSTEM SHOWS A 250 HR BREAK-IN PERIOD FOLLOWED BY STABILIZATION

Raw data analysis from long term testing

- Cell 3 shows R_{Ω} instability, likely electrical contact issue (can be corrected for)
- Cell 2 change in performance due to gas composition (OCV) change
- Furnace controller issue at 375hr

Durability trends identified

- LNO stabilizing after 250 hrs
- Cells 1, 2, 3 show reproducible behavior (cell manuf. and testing reliability)
- Cell 4 shows stability issues, unclear if internal or external



Collaboration: Effectiveness

Collaboration with EMN project node experts

- Monthly conference calls with the full team including EMN project node experts
- Discussions with team at INL (James O'Brian, Dong Ding, Hanping Ding) refined post-testing analysis plan including CT scanning of button cells
- Discussions with Eric Coker at SNL focused on quantification of oxygen hyperstoichiometry. Through this discussion atmosphere controlled TGA was added to the project plan to supplement atmosphere controlled XRD. In this way the risk of being able to measure this property was reduced
- Data analysis discussions after significant testing milestones were used to share progress, interpret results, and plan next steps



PROJECT TASKS WERE SPREAD ACROSS THE TEAMS, TAKING FULL ADVANTAGE OF BOTH EXPERTISE AND RELEVANT UNIQUE CAPABILITIES



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Next Steps

- Button cell fabrication and testing of top candidates
- Investigate cross-family stability and performance
- Engage with additional Nodes for modeling support
- Additional focus on potential interfacial reactions
- Optimization of stoichiometry and sintering to maximize performance and minimize degradation
- Down selection of electrode compositions for integration into short stacks
- Initial short stack performance and durability testing



HydroGEN: Advanced Water Splitting Materials







Any proposed future work is subject to change based on funding levels 24



Performance of button cells with a Nickelate oxygen electrode

- Dramatic performance increase
 - 70% higher electrolyzer current density @ 1.4V than LSM-YSZ
 - Indicates Nickelate ceria interactions prevented during sintering
- Interface stable/well-adhered after 336 hr of continuous operation



Status vs. 3-year targets

Metric	Target	Current
ASR	<u><</u> 0.3 ohm cm²	0.24 ohm cm^2
Current Density	<u>></u> 1 A/cm ² @ 1.4V	1.2 A/cm² @ 1.4V
Degradation Rate	<u><</u> 0.3 %/khr	LSM > LNO >> 0.3%

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- SG: S. Soulekar, R. Wang

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Technical Backup Slides (optional)

Accomplishments

Summary of milestones and progress

Milestone	Description	Month	Metric	Progress
M1.1	Powders meet specifications	3	Stoichiometry met (ICP)Phase purity (XRD)	 √ Process defined √ Stoichiometry and purity confirmed √ NNO, NDC , LNO, LDC, complete
M1.2	Identification of stability boundaries	8	Decomposition free regions determined for nickelate-ceria mixtures	• √ Completed
M1.3	Determine top 3 composites	12	 >96 S/cm @800C Oxygen exchange coefficient 1.2x10⁻⁵ cm/s Maps of oxygen nonstoichiometry f(T, PO2) 	• √ Measurements completed
M2.1	Button cell fabrication	5	 Baseline button cells co-sintered with He leak (in air) <1x10-8 mbarLs⁻¹cm⁻² Microstructure acceptable 	 √ Baseline cells prepared √ Initial characterization started √ Tapes of nickelate prepared √ Nickelate microstructure to be improved
M2.1	Accelerated testing	11	 Baseline degradation rate established in accelerated testing Degradation mode identified microstructurally 	 √ Baseline cells tested √ Baseline cells examined √ Nickelate cells to be tested Nickelate cells to be examined
G/NG	Button cell test results operating in electrolysis conditions	12	 25% higher electrolyzer current density @ 1.4V than baseline LSM-YSZ Well adhered layers after 2 wks of operation Degradation rate <1.5%/1000 hr 	Discussion

Accomplishments: Stability Boundaries

STABLE REGIONS HAVE BEEN IDENTIFIED FOR FURTHER WORK, IN THESE DOMAINS THE REACTION BETWEEN NICKELATE AND CERIA PHASES IS MINIMIZED OR ELIMINATED. THIS PREVENTS LOSS OF THE ACTIVE NICKELATE PHASE AND IMINATION OF RESISTIVE GRAIN BOUNDARY PHASES.

Electrocatalyst	Ionic Conductor	Stability of System
La ₂ NiO ₄ (LNO)	La-doped CeO ₂ (LDC)	\checkmark
Sm_2NiO_4 (SNO)	Sm-doped CeO ₂ (SDC)	×
Nd ₂ NiO ₄ (NNO)	Nd-doped CeO ₂ (NDC)	\checkmark

- Identified that a relatively high level of • doping is required to prevent ceria from decomposing the nickelate phase
- Was not able to effectively utilize Sm ۲ during Phase 1 testing
- Did not investigate cross family stabilization - potential Phase 2 investigation



Ceria Phase Lattice Parameter

5.575

Initial OCVs are close to predicted values

Accomplishments: Good Cell Integrity

CTE DIFFERENCES DID NOT MANIFEST AS ELECTROLYTE CRACKS, THIS WAS A CRITICAL RESULT IN TERMS OF ENABLING CO-SINTERING WHICH RESULTS IN WELL BONDED INTERFACES THROUGHOUT THE CELL



OCV is stable over time